Approved for use through 07/31/2006. OMB 0651-0031 U.S. Patent and Trademark Office: U.S. DEPARTMENT OF COMMERCE

Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it contains a valid OMB control number.

ADENSE Citute for form 1449A/PTO

Sheet

INFORMATION DISCLOSURE STATEMENT BY APPLICANT

(use as many sneets	as necessar	<i>y</i> /
1	of	6

Complete if Known					
Application Number	10/602,693				
Filing Date	June 20, 2003				
First Named Inventor	Sommadossi et al.				
Group Art Unit	1614				
Examiner Name	Unassigned				
Attorney Docket Number	06171.105070 IDX 1006 DIV				

3425621_1

	U.S. PATENT DOCUMENTS							
Examiner Initials *	Cite No. 1		ment Kind Code if known)	Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pgs, Clmns, Lns, Where Relevant Passages/Relevant Figs Appear		
B	AA_	3,480,613	Α	Walton et al.	11-25-1969			
	AB	5,977,061	Α	De Clercq	11-02-1999			
	AC	6,340,690	Bl	Bachand et al.	01-22-2002			
	AD	6,348,587	BI	Schinazi et al.	02-2002	•		
	AE	6,395,716	Bl	Gosselin et al. (Novirio / Idenix)	05-28-2002			
	AF	6,444,652	BI	Gosselin et al. (Novirio / Idenix)	09-03-2002			
	AG	6,573,248	Bl	Ramasamy et al.	06-03-2003			
	AH	2002/0019363	Al	Ismaili et al.	02-2002			
	AI	2002/0055483	Al	Watanabe et al.	05-09-2002			
	AJ	2002/0147160	Al	Bhat et al.	10-10-2002			
	AK	2003/008841	Al	Devos et al.	01-09-2003			
	AL	2003/028013	Al	Wang et al.	02-06-2003			
	AM	2003/0050229	Al	Sommadossi et al.	03-13-2003			
	AN	2003/0060400	Al	LaColla et al.	03-27-2003			
JK.	AO	2003/0083307	Al	Devos et al.	05-01-2003			
10/	AP	2003/0087873	Al	Stuyver et al.	05-08-2003			

				FOR	EIGN PATENT DOCUMENTS			
Examiner Initials *	Cite No. 1		ign Patent Docum Number Kind (if k		Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD- YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	I _e
(le	AQ	FR	1,521,076	Α	Merck & Co. Inc.	04-12-1968		
	AR	FR	1,581,628	Α	Merck & Co. Inc.	09-19-1969		
	AS	FR	2,662,165	Α	Univ. Paris Curie	11-22-1991		
	AT	GB	1,163,103	Α	Merck & Co. Inc.	09-04-1969		
	ΑŬ	GB	1,209,654	Α	Merck & Co. Inc.	10-21-1970		
	AV	JP	63-215694	Α	Yamasa Shoyu Co. Ltd.	09-08-1988		
	AW	JP	06-228186	Α	Yamasa Shoyu Co. Ltd.	08-16-1994		
	AX	WO	98/16184	A2	ICN Pharmaceuticals	04-23-1998		
	AY	WO	99/43691	A1	Emory U.; U.Ga.R.F.	02-09-1999		
Y)	AZ	WO	00/09531	A2	Novirio Pharm. (Idenix)	02-24-2000		
	AAA	WO	01/32153	A2	Biochem Pharma	05-10-2001		

Examiner	Date Considered	3/29/05
Signature	Considered	1/2 1/05

^{*}EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

Burden Hour Statement: This form is estimated to take 2.0 hours to complete. Time will vary depending upon the needs of the individual case. Any comments on the amount of time you are required to complete this form should be sent to the Chief Information Officer, U. S. Patent and Trademark Office, Washington, DC 20231. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Assistant Commissioner for Patents, Washington, DC 20231.

¹ Unique citation designation number. ² See attached Kinds of U.S. Patent Documents. ³ Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3). ⁴ For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. 5 Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST. 16 if possible. Applicant is to place a check mark here if English language Translation is attached.

Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it contains a valid OMB control number. Complete if Known Substitute for form 1449A/PTO **Application Number** 10/602,693 Filing Date June 20, 2003 INFORMATION DISCLOSURE First Named Inventor STATEMENT BY APPLICANT Sommadossi et al. **Group Art Unit** 1614 Examiner Name Unassigned (use as many sheets as necessary) **Attorney Docket Number** 06171.105070 IDX 1006 DIV of 6 Sheet 2

3425621 1

				FOR	EIGN PATENT DOCUMENTS			
Examiner Initials *	Cite No. 1		ign Patent Docum Number Kind ((if ki		Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD- YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T6
N	BA	WO	01/60315	A2_	Biochem Pharma	08-23-2001		
	BB	WO	01/68663	Al	ICN Pharmaceuticals	09-20-2001		
	BC	WO	01/79246	A2	Pharmasset	10-25-2001		
	BD	WO	01/90121	A2	Novirio Pharm. (Idenix)	11-29-2001		
	BE	WO	01/91737	A2	Novirio Pharm. (Idenix)	06-12-2001	•	
	BF	WO	01/92282	A2	Novirio Pharm. (Idenix)	06-12-2001		
	BG	WO	01/96353	A2	Novirio Pharm. (Idenix)	12-20-2001		
	BH	WO	02/03997	Al	ICN Pharmaceuticals	01-17-2002		
	BI	WO	02/18404	A2	F. Hoffmann-La Roche	03-07-2002		
	BJ	WO	02/32920	A2	Pharmasset	04-25-2002		
	BK	wo	02/48165	A2	Pharmasset	06-20-2002	-	
	BL	wo	02/057287	A2	Merck & Co. Inc.	07-25-2002	-	
	BM	WO	02/057425	A2	Merck & Co. Inc.	07-25-2002		
	BN	WO	02/070533	A2	Pharmasset	09-12-2002		
	ВО	wo	02/094289	A1	F. Hoffmann-La Roche	11-28-2002		
	BP	WO	02/100415	A2	F. Hoffmann-La Roche	12-19-2002		
	BQ	wo	03/026589	A2	Idenix; CNRS; U. Montp.	04-03-2003		
	BR	wo	03/026675	A 1	Idenix; CNRS; U. Montp.	04-03-2003		
	BS	wo	03/051899	A 1	Ribapharm	06-26-2003		
	BT	wo	03/061385	A 1	Ribapharm	07-31-2003		
	BU	WO	03/061576	A2	Ribapharm	07-31-2003		
	BV	wo	03/062255	A2	Ribapharm	07-31-2003		1
	BW	wo	03/062256	A1	Ribapharm	07-31-2003		
	BX	WO	03/062257	A1	Ribapharm	07-31-2003		
	BY	WO	03/063771	A2	Pharmasset	08-07-2003		
	BZ	WO	03/068162	A2	Pharmasset	08-21-2003		
	BAA	wo	03/072757	A2	Biota Inc.	09-04-2003		
1//	BAB	WO	03/093290	A2	Genelabs Technologies	11-13-2003		
	BAC	wo	04/002422	A2	Idenix; Univ.D.S.Cagliari	01-08-2004		
(4)	BAD	wo	04/002999	A2	Idenix; Univ.D.S.Cagliari	01-08-2004		

Examiner		Date	2/2/-
Signature	1	Considered	3/29/05

^{*}EX AMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

¹Unique citation designation number. ²See attached Kinds of U.S. Patent Documents. ³Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3). ⁴For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. ⁵Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST. 16 if possible. ⁶Applicant is to place a check mark here if English language Translation is attached.

Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it contains a valid OMB control number

Cubatha	Substitute for form 1440 A POTO			Complete if Known		
Substitute for form 1449A/PTO				Application Number	10/602,693	
INFO	RMATION I	DISCLO	SURE	Filing Date	June 20, 2003	
STATEMENT BY APPLICANT		First Named Inventor	Sommadossi et al.			
				Group Art Unit	1614	
(use as many sheets as necessary)		Examiner Name	Unassigned			
Sheet	3	of	6	Attorney Docket Number	06171.105070 IDX 1006 DIV	

3425621 1 OTHER PRIOR ART - NON PATENT LITERATURE DOCUMENTS Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, Examiner Cite No. 1 journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published. Tڻ Initials • ALTMANN et al, "The synthesis of 1'-methyl carbocyclic thymidine and its effect on nucleic acid duplex stability," Synlett, Thieme Verlag, Stuttgart, De, 10:853-855 (1994). BAGINSKI, S. G, et al., "Mechanism of action of a pestivirus antiviral compound," PNAS USA, 97(14):7981-7986 (2000). BEIGELMAN, L.N., et al, "Epimerization during the acetolysis of 3-O-acetyl-5-O-benzoyl-1,2-O-CC isopropylidene-3-C-methyl- α ,D-ribofuranose. Synthesis of 3'-C-methylnucleosides with the β -Dribo- and α-D-arabino configurations," Carbohydrate Research, 181:77-88 (1988). BEIGELMAN, L.N., et al, "A general method for synthesis of 3'-C-alkylnucleosides," Nucleic Acids CD Symp. Ser., 9:115-118 (1981). BERENGUER, M., et al, "Hepatitis B and C viruses: Molecular identification and targeted antiviral CE therapies," Proceedings of the Association of American Physicians, 110(2), 98-112 (1998). CARROLL, S.S., et al., "Inhibition of hepatitis C virus RNA replication by 2'-modified nucleoside analogs," The Journal of Biological Chemistry, 278(14):11979-11984 (2003). CZERNECKI, S., et al, "Synthesis of various 3'-branched 2',3'-unsaturated pyrimidine nucleosides as CG potential anti-HIV agents," J. Org. Chem., 57:7325-7328 (1992). De FRANCESCO, R., et al., "Approaching a new era for hepatitis C virus therapy: Inhibitors of the CH NS3-4A serine protease and the NS5B RNA-dependent RNA polymerase," Antiviral Research, 58:1-16 (2003). FAIVRE-BUET, V., et al, "Synthesis of 1'-deoxypsicofuranosyl-deoxynucleosides as potential anti-HIV agents," Nucleosides & Nucleotides, 11(7):1411-1424 (1992). FARKAS, J., et al., "Nucleic acid components and their analogues. XCIV. Synthesis of 6-amino-9-(1deoxy-β-D-psicofuranosyl)purine", Collect. Czech. Chem. Commun. 32:2663-2667 (1967). FARKAS, J., et al., "Nucleic acid components and their analogues. LXXIX. Synthesis of methyl 1 deoxy-D-psicofuranosides substituted at C(1) with halo atoms or a mercapto group," Collect. Czech. Chem. Commun., 31:1535-1543 (1996). FEDOROV, I.I., et al, "3'-C-Branched 2'-deoxy-5-methyluridines: Synthesis, enzyme inhibition, and antiviral properties," J. Med. Chem., 35(24):4567-4575 (1992). FRANCHETTI, P., et al., "2'-C-Methyl analogues of selective adenosine receptor agonists: synthesis **CM** and binding studies," J. Med. Chem., 41(10):1708-1715 (1998). GROUILLER, A., et al., "Novel p-toluenesulfonylation and thionocarbonylation of unprotected CN thymine nucleosides," Synlett, 1993, 221-222 (March 1993). HARAGUCHI, K., et al., "Preparation and reactions of 2'- and 3'- vinyl bromides of uracil CO nucleosides: Versatile synthons for anti-HIV agents," Tetrahedron Letters, 32(28):3391-3394 (1991).

Examiner Signature	/	Date Considered	3/27/05

^{*}EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

¹Unique citation designation number. ² See attached Kinds of U.S. Patent Documents. ³ Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3). ⁴ For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. ⁵ Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST. 16 if possible. ⁶ Applicant is to place a check mark here if English language Translation is attached.

3425621 1

U.S. Patent and Trademark Office: U.S. DEPARTMENT OF COMMERCE

_	Under the Paperwon	k Reduction	Act of 1993, its person	is the required to respond to a confection	in of information unless it contains a valid OMLB control number.	
Cubetine	for form 1449A/PTO			Complete if Known		
Susanu	e for form 1449A/PTO			Application Number	10/602,693	
INFO	DRMATION D	ISCL	OSURE	Filing Date	June 20, 2003	
STA	TEMENT BY	APPLI	CANT	First Named Inventor	Sommadossi et al.	
				Group Art Unit	1614	
	(use as many sheets	as necessai	(ער	Examiner Name	Unassigned	
Sheet	4	of	6	Attorney Docket Number	06171.105070 IDX 1006 DIV	

OTHER PRIOR ART - NON PATENT LITERATURE DOCUMENTS Cite Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, Examiner No. 1 journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published. Initials * HARAGUCHI, K., et al., "Stereoselective synthesis of 1'-C-branched uracil nucleosides from DA uridine," Nucleosides & Nucleotides, 14(3-5):417-420 (1995). HARRY-O'KURU, R.E., et al., "A short, flexible route toward 2'-C-branched ribonucleosides", DB J.Org. Chem., 62:1754-1759 (1997). (Scheme 11). DC HARRY-O'KURU, R.E., et al., "2'-C-Alkylribonucleosides: Design, synthesis, and conformation," Nucleosides & Nucleotides, 16(7-9):1457-1460 (1997). ["Rogers" in #2; correct name in #7] HATTORI, H., et al, "Nucleosides and nucleotides. 175. Structural requirements of the sugar moiety DD for the antitumor activities of new nucleoside antimetabolites, 1-(3-C-ethynyl-b-D-ribopentofuranosyl)cytosine and -uracil," J. Med. Chem., 41:2892-2902 (1998). HREBABECKY, H., et al., "Nucleic acid components and their analogues. CXLIX. Synthesis of pyrimidine nucleosides derived from 1-deoxy-D-psicose," Collect. Czech. Chem. Commun., 37:2059-2065 (1972). HREBABECKY, H., et al. "Synthesis of 7- and 9β-D-psicofuranosylguanine and their 1'-deoxy DF derivatives," Collect. Czech. Chem. Commun., 39:2115-2123 (1974). IINO, T., et al., "Nucleosides and nucleotides. 139. Stereoselective synthesis of (2'S)-2'-C-alkyl-2'-DG deoxyuridines," Nucleosides and Nucleotides, 15(1-3):169-181 (1996). ITOH, Y., et al, "Divergent and stereocontrolled approach to the synthesis of uracil nucleosides DH branched at the anomeric position," J. Org. Chem., 60(3):656-662 (1995). JOHNSON, C.R., et al, "3'-C-Trifluoromethyl ribonucleosides," Nucleosides & Nucleotides, DI 14(1&2):185-194 (1995). KAWANA, M., et al., "The deoxygenation of tosylated adenosine derivatives with Grignard DJ reagents," Nucleic Acids Symp. Ser., 17:37-40 (1986). LAVAIRE, S., et al., "3'-Deoxy-3'-C-trifluoromethyl nucleosides: Synthesis and antiviral DK evaluation," Nucleosides & Nucleotides, 17(12):2267-2280 (1998). LEYSSEN, P. et al., "Perspectives for the treatment of infections with Flaviviridae," Clinical Microbiology Reviews (Washington, D.C.), 13(1):67-82 (January 2000). MARTIN, X., et al., "Intramolecular hydrogen bonding in primary hydroxyl of thymine 1-(1-deoxy-\beta-DM D-psicofuranosyl) nucleoside," Tetrahedron, 50(22):6689-6694 (1994). MATSUDA, A., et al., "Radical deoxygenation of tert-alcohols in 2'-branched-chain sugar pyrimidine DN nucleosides: Synthesis and antileukemic activity of 2'-deoxy-2'(S)-methylcytidine," Chem. Pharm. Bull., 35(9):3967-3970 (1987). MATSUDA, A., et al., "Alkyl addition reaction of pyrimidine 2'-ketonucleosides: Synthesis of 2'-DO branched-chain sugar pyrimidine nucleosides (Nucleosides and Nucleotides. LXXXI)," Chem. Pharm. Bull., 36(3):945-953 (1988).

Examiner Signature	1 Camero	Date Considered	3/29/05	

^{*}EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

Unique citation designation number. ² See attached Kinds of U.S. Patent Documents. ³ Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3). ⁴ For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. ⁵ Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST. 16 if possible. ⁶ Applicant is to place a check mark here if English language Translation is attached.

Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it contains a valid OMB control number. Complete if Known Substitute for form 1449A/PTO Application Number 10/602,693 Filing Date INFORMATION DISCLOSURE June 20, 2003 First Named Inventor STATEMENT BY APPLICANT Sommadossi et al. Group Art Unit 1614 Examiner Name Unassigned (use as many sheets as necessary) **Attorney Docket Number** 06171.105070 IDX 1006 DIV 6 5 of Sheet

3425621 1

		OTHER PRIOR ART – NON PATENT LITERATURE DOCUMENTS	
Examiner	Cite	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine,	
Initials *	No.	journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	Té
	EA	MATSUDA, A., et al., "Nucleosides and Nucleotides. 94. Radical deoxygenation of tert-alcohols in	
(4)		1-(2-C-alkylpentofuranosyl)pyrimidines: Synthesis of (2'S)-2'-deoxy-2'-C-methylcytidine, an	
(V)		antileukemic nucleoside, " J. Med. Chem., 34:234-239 (1991).	
1	EB	MATSUDA, A., et al., "Nucleosides and Nucleotides. 104. Radical and palladium-catalyzed	
		deoxygenation of the allylic alcohol systems in the sugar moiety of pyrimidine nucleosides,"	
		Nucleosides & Nucleotides, 11(2/4):197-226 (1992).	
	EC	MIKHAILOV, S.N., et al., "Synthesis and properties of 3'C-methylnucleosides and their phosphoric	
		esters," Carbohydrate Research, 124:75-96 (1983).	
	ED	MIKHAILOV, S.N., et al., "Substrate properties of C'-methylnucleoside and C'-methyl-2'-	
		deoxynucleoside 5'-triphosphates in RNA and DNA synthesis reactions catalysed by RNA and DNA	•
		polymerases," Nucleosides & Nucleotides, 10(1-3):339-343 (1991).	
	EE	MIKHAILOV, S.N., et al, "Hydrolysis of 2'- and 3'-C-methyluridine 2'c3'-cyclic monophosphates	
		and interconversion and dephosphorylation of the resulting 2'- and 3'-monophosphates: Comparison	
1 1		with the reactions of uridine monophosphates," J. Org. Chem., 57 (15):4122-4126 (1992).	
	EF	NUTT, R.F., et al., "Branched-chain sugar nucleosides. III. 3'-C-methyladenine", J. Org. Chem.,	
		33:1789-1795 (1968).	
	EG	OIVANEN, M., et al, "Additional evidence for the exceptional mechanism of the acid-catalyzed	
		hydrolysis of 4-oxopyrimidine nucleosides: Hydrolysis of 1-(1-alkoxyalkyl)uracils, seconucleosides,	
		3'-C-alkyl nucleosides and nucleoside 3',5'-cyclic monophosphates," J. Chem. Soc. Perkin Trans. 2,	
		1994:309-314 (1994).	
	EH	ONG, S.P., et al, "Synthesis of 3'-C-methyladenosine and 3'-C-methyluridine diphosphates and their	
		interaction with the ribonucleoside diphosphate reductase from Corynebacterium nephridii,"	
		Biochemistry, 31(45):11210-11215 (1992).	
	El	Oral Session V, Hepatitis C Virus, Flaviviridae; 16th International Conference on Antiviral Research	
		(April 27, 2003, Savannah, Ga.) p A75-77.	
	EJ	PAN-ZHOU, X-R, et al., "Differential effects of antiretroviral nucleoside analogs on mitochondrial	
		function in HepG2 cells," Antimicrob. Agents Chemother., 44:496-503 (2000).	
	EK	ROSENTHAL, A., et al., "Branched-chain sugar nucleosides. Synthesis of 3'-C-ethyl (and 3'-C-	
レン	· -	butyl)uridine Carbohydrate Research, 79:235-242 (1980).	
	EL	SAMANO, V., et al., "Synthesis and radical-induced ring-opening reactions of 2'-deoxyadenosine-2'-	
(4)		spirocyclopropane and its uridine analogue. Mechanistic probe for ribonucleotide reductases," J. Am.	
		Chem. Soc., 114:4007-4008 (1992).	
		Chem. 200., 117.7001 7000 (1772).	1

Examiner Signature	Date Considered	3/29/05	

^{*}EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

¹Unique citation designation number. ²See attached Kinds of U.S. Patent Documents. ³Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3). ⁴ For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. ⁵ Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST. 16 if possible. ⁶ Applicant is to place a check mark here if English language Translation is attached.

Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it contains a valid OMB control number.

				Complete if Known		
Substitute for form 1449A/PTO				Application Number	10/602,693	
INFORMATION DISCLOSURE STATEMENT BY APPLICANT			SURE	Filing Date	June 20, 2003	
			CANT	First Named Inventor	Sommadossi et al.	
				Group Art Unit	1614	
	(use as many sheets as necessary)			Examiner Name	Unassigned	
Sheet	6	of	6	Attorney Docket Number	06171.105070 IDX 1006 DIV	

3425621 1

•		OTHER PRIOR ART – NON PATENT LITERATURE DOCUMENTS	
Examiner	Cite	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine,	T
Initials *	No. 1	journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published. SAMANO, V., et al., "Nucleic acid related compounds. 77. 2',3'-Didehydro-2',3'-dideoxy-2'(and	
	FA		f
(1,)		3')-methylnucleosides via [3,3]-sigmatropic rearrangements of 2'(and 3')-methylene-3'(and 2')-O-thiocarbonyl derivatives and radical reduction of a 2'-chloro-3'-methylene analogue," Can. J. Chem.,	
(17)		•	
	ED	71:186-191 (1993).	-
	FB	SCHMIT, C., et al, "The effects of 2'- and 3'-alkyl substituents on oligonucleotide hybridization and	
		stability," Biorganic & Medicinal Chemistry Letters, 4(16):1969-1974 (1994). ["Altmann"]	-
	FC	SERAFINOWSKI, P.J., et al., "New method for the preparation of some 2'- and 3'-trifluoromethyl-	
		2',3'-dideoxyuridine derivatives," <i>Tetrahedron</i> (Elsevier Science Publishers), 56(2):333-339 (1999).	-
	FD	SHARMA, P.K., et al., "Synthesis of 3'-trifluoromethyl nucleosides as potential antiviral agents,"	
	- DD	Nucleosides, Nucleotides and Nucleic Acids, 19(4):757-774 (2000).	
	FE	SOMMADOSSI J-P, et al., "Comparison of cytotoxicity of the (-)- and (+)-enantiomer of 2',3'-	
		dideoxy-3'-thiacytidine in normal human bone marrow progenitor cells" Biochemical Pharmacology,	
	FF	44:1921-1925 (1992).	
	FF	SOMMADOSSI J-P, et al., "Toxicity of 3'-azido-3'-deoxythymidine and 9-(1,3-dihydroxy-2-propoxymethyl)guanine for normal human hematopoietic progenitor cells in vitro" Antimicrobial	•
		Agents and Chemotherapy, 31:452-454 (1987).	
	FC		
	FG	TRITSCH, D., et al., "3'-β-ethynyl and 2'-deoxy-3'-β-ethynyl adenosines: First 3'-β-branched adenosines substrates of adenosine deaminase," Bioorganic & Medicinal Chemistry Letters, 10:139-	
		141 (2000).	
	FH	TUNITSKAYA, V.L., et al., "Substrate properties of C'-methyl UTP derivatives in T7 RNA	
	rn.	polymerase reactions. Evidence for N-type NTP conformation," FEBS Letters, 400:263-266 (1997).	
	FI	USUI, H., et al., "Synthesis of 2'-deoxy-8,2'-ethanoadenosine and 3'-deoxy-8,3'-ethanoadenosine	
	r.i	(Nucleosides and Nucleotides. LXIV)," Chem. Pharm. Bull., 34(1):15-23 (1986).	
	FJ	WALCZAK, K., et al., "Synthesis of 1-(3-alkyl-2,3-dideoxy-D-pentofuranosyl)uracils with potential	
		anti-HIV activity," Acta Chemica Scand., 45:930-934 (1991).	
	FK	WALTON, E., et al., "Branched-chain sugar nucleosides. V. Synthesis and antiviral properties of	
		several branched-chain sugar nucleotides," J. Med. Chem., 12:306-309 (1969).	
17	FL	WOLFE, M.S., et al., "A concise synthesis of 2'-C-methylribonucleosides," Tetrahedron Letters,	
		36(42):7611-7614 (1995).	
	FM	WU, JC., et al., "A new stereospecific synthesis of [3.1.0] bicyclic cyclopropano analog of 2',3'-	
17	,	dideoxyuridine, Tetrahedron, 46(7):2587-2592 (1990).	
			_

Examiner Signature	Date Considered	3/29/05	
Signature	Considered	7/2//03	

^{*}EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

¹ Unique citation designation number. ² See attached Kinds of U.S. Patent Documents. ³ Enter Office that issued the document, by the two-letter code (WIPO Standard ST.3). ⁴ For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. ⁵ Kind of document by the appropriate symbols as indicated on the document under WIPO Standard ST. 16 if possible. ⁶ Applicant is to place a check mark here if English language Translation is attached.